



(11) **EP 1 110 691 A2**

(12) **EUROPEAN PATENT APPLICATION**

(43) Date of publication:
27.06.2001 Bulletin 2001/26

(51) Int Cl.7: **B29B 7/42**, B29B 7/84,
B29B 7/88, B29K 83/00,
C08J 3/20

(21) Application number: 00310967.5

(22) Date of filing: 08.12.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR
Designated Extension States:
AL LT LV MK RO SI

- Giammattei, Mark Howard
Selkirk, New York 12158 (US)
- Carbone, John William
Schenectady, New York 12309 (US)
- Singh, Navjot
Clifton Park, New York 12065 (US)
- Rocha, Gerardo
Waterford, New York 12188 (US)
- Banevicius, John Peter
Clifton Park, New York 12065 (US)

(30) Priority: 21.12.1999 US 468622

(71) Applicant: **GENERAL ELECTRIC COMPANY**
Schenectady, NY 12345 (US)

(72) Inventors:
• Silvi, Norberto
Clifton Park, New York 12065 (US)
• Dean, Robert Edward
Rexford, New York 12148 (US)

(74) Representative: Szary, Anne Catherine, Dr. et al
London Patent Operation,
GE International, Inc.,
Essex House,
12-13 Essex Street
London WC2R 3AA (GB)

(54) **Compounding filled silicone compositions**

(57) High levels of inorganic filler, processing fluid and silicone polymer are compounded and devolatilized into homogeneous filled compositions with requisite reinforcing properties and levels of volatiles. In the method, a filler, processing fluid and silicone polymer are

mixed in a first compounding apparatus (12,44,76) to produce a dispersed composition containing volatiles. The dispersed composition is then compounded in a long extruder (14,46,78) having an L/D ratio of greater than 50 to devolatilize the dispersed composition.

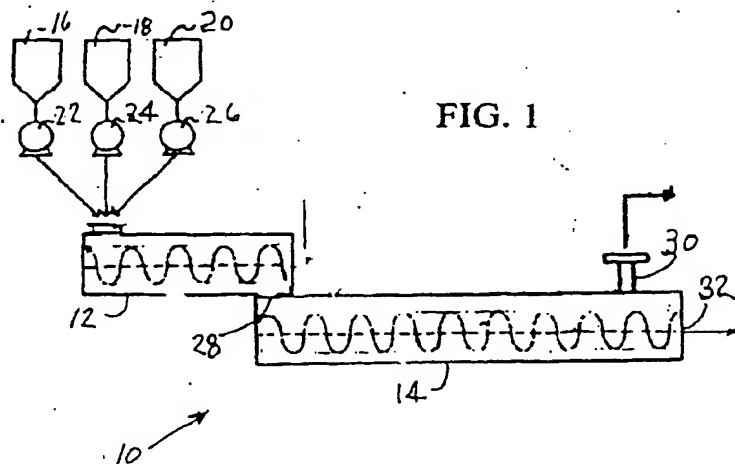


FIG. 1

Description

[0001] The invention relates to a method of compounding heat-vulcanizable silicone compositions.

[0002] A heat-vulcanizable silicone composition comprises a high viscosity silicone polymer, an inorganic reinforcing filler and various additives that aid processing or impart desired final properties to the composition. A vulcanizing agent can be added and the composition heat-cured to fabricate silicone rubber moldings such as gaskets, medical tubing and computer keypads.

[0003] Typically, the heat-vulcanizable silicone composition is produced by kneading a high-viscosity polydiorganosiloxane, the inorganic filler and additives by means of a batch kneading machine such as a high intensity Banbury mixer or a low intensity double arm dough mixer. In this process, polydiorganosiloxane, inorganic filler and treating agents are batch mixed until desired properties are obtained. This process requires long residence times and large amounts of energy. Non-homogeneous shear and extensional stresses across a commercial sized batch can result in non-uniform size distribution of filler that results in variations in properties. Batches processed at different times may be characterized by different physical properties. The batch process is labor, energy and capital intensive and produces materials of only marginal consistency.

[0004] In Kasahara *et al.*, U.S. Pat. 5,198,171, a pre-concentrate of polydiorganosiloxane, inorganic filler and treating agents is formed by a high speed mechanical shearing mixer. The resulting premix is further compounded in a same-direction double screw extruder. The premix is formed in a first step wherein a diorganopolysiloxane having a viscosity at 25°C of 1×10^5 cP or more, an inorganic filler and a treating agent are mixed in a high-speed mechanical shearing machine to provide a flowable particulate mixture in which each ingredient is present in a substantially uniform, finely dispersed state. The flowable particulate mixture is then fed at a constant feed rate into a kneading and extruding machine that has two screws rotating in the same direction and has a length to diameter ratio (L/D) of 25 to 50.

[0005] In Hamada *et al.*, U.S. Pat. 5,409,978, a pre-concentrate of polydiorganosiloxane, inorganic filler and treating agents is formed at a temperature in the range of about 200°C to 300°C in a co-rotating continuous double screw extruder with L/D of about 25 to 50. The pre-concentrate is then compounded and heat treated at 150°C to 300°C in a counter-rotating, double screw extruder. A useful L/D ratio for the extruder is in the range of about 5 to 15.

[0006] Highly vigorous first step processes for forming a composition of silicone polymer, filler and treating agent generate a product having a high volatiles content. The composition must be devolatilized in a second step to produce a useful product. Devolatilization is important for at least two reasons. First, devolatilization removes unreacted treating agents to "quench" further reaction with filler silanol groups. Otherwise, further reaction can diminish filler reinforcement properties in the heat-vulcanizable silicone composition. Second, devolatilization removes surplus liquids that adversely affect rheological properties of the silicone composition. Residual volatiles in the filled silicone composition should not exceed the level of about 2% by weight of the final formulation. Preferably, residual volatiles in the filled silicone composition should not exceed the level of about 1%.

[0007] However, devolatilization is not complete in known two-step processes. There is a need for a process that continuously and consistently produces a devolatilized high viscosity filled silicone polymer composition from filler, additive and polymer. Further, there is a need for an improved continuous compounding process.

BRIEF SUMMARY OF THE INVENTION

[0008] The invention provides a process that compounds high levels of inorganic filler, processing fluid and silicone polymer into homogeneous filled and devolatilized silicone compositions with requisite reinforcing properties and levels of volatiles. The method comprises mixing a filler, processing fluid and silicone polymer into a first compounding apparatus to produce a dispersed composition containing volatiles. The dispersed composition is then compounded in an extruder having an L/D ratio of greater than 50 to devolatilize the dispersed composition.

[0009] In another aspect, a process of compounding a filled silicone composition comprises forming a premix of filler and silicone polymer in a first mixer. The premix is then compounded with further filler and silicone polymer and added processing fluid in a first compounding apparatus to produce a dispersed composition containing volatiles. The dispersed composition is devolatilized in an extruder having an L/D ratio of greater than 50.

[0010] In still another aspect, a system is provided for compounding filled silicone compositions. The system comprises a first compounding apparatus and an extruder having an L/D ratio of greater than 50. The extruder is connected to the first compounding apparatus to continuously receive a dispersed composition containing volatiles.

BRIEF DESCRIPTION OF THE DRAWINGS

[0011]

FIG. 1 is a schematic representation of a silicone composition compounding process and system;

FIG. 2 is a schematic representation of another silicone composition compounding process and system; and

FIG. 3 is a schematic representation of still another silicone composition compounding process and system.

DETAILED DESCRIPTION OF THE INVENTION

[0012] A process and system are provided by the invention to compound high levels of components such as treated fumed silica and processing fluid into a silicone polymer, such as high molecular weight poly(dimethylsiloxane). The process and system produce a homogeneous mixture having required reinforcing properties and levels of residual volatiles. The process includes at least two continuous compounding steps, wherein filler and silicone polymer are mixed and compounded in a first extruder and devolatilized in a long second extruder. A long extruder is defined as an extruder having a length to diameter ratio of at least about 50. In one embodiment, the long extruder can have a length to diameter ratio of at least about 60.

[0013] The processing fluid of the invention is a fluid that can be admixed with a filler and compounded to density a filler for further processing. The processing fluid can also provide a processing function. It can be a liquid treating agent, plasticizer, flow improving additive, cross-linking agent, water or inert blanketing gas. Silicone polymers of a molecular weight greater than 7000 are not processing fluids. Preferably, the processing fluid is a liquid treating agent such as a silanol-reacting treating agent that can be added before, with or after addition of filler to wet the filler to reduce overall processing time for reaction between functional groups in the treating agent and silanols on the surface of the filler.

[0014] In one embodiment, the processing fluid is a solution prepared by mixing (in weight) 1.21 parts of a silanol-stopped polydimethylsiloxane, 1.82 parts of a vinyl-stopped dimethyl-methylvinylsiloxane and 0.12 part of a hydroxy-terminated polydimethyl-methylvinylsiloxane. A broad range of parts of silanol-stopped polydimethylsiloxane/vinyl-stopped dimethyl-methylvinylsiloxane/hydroxy-terminated polydimethyl-methylsiloxane can be 0.49/0.73/0.05 to 1.93/2.91/0.19, a desired range is 0.85/1.27/0.08 to 1.57/2.37/0.16 and a preferred range is 1.09/1.64/0.11 to 1.32/2.0/0.13.

[0015] In another embodiment, the processing fluid can be a combination of treating agent, preferably HMDZ and water. This combination can comprise a weight ratio of treating agent/water of between about 0.05 to about 50 or between about 0.1 and about 20 or between about 1 and about 6. The HMDZ can be added either together with a premix or separately at a single or at multiple locations of an extruder or to a batch mixer in the compounding step.

[0016] The processing fluid can be combined with filler in a weight proportion of about 0.1 to about 100 parts fluid to 100 parts of filler, desirably about 0.5 to about 75 parts fluid to 100 parts of filler and preferably about 1.0 to about 50 parts fluid to 100 parts of filler. The processing fluid can be added at a single location or at a plurality of locations for step treatment of the filler.

[0017] The inorganic filler that can be used in the invention can be any inorganic filler used in blends with silicone polymers. Examples of inorganic fillers include a reinforcing silica such as fumed silica or precipitated silica or a silica that has been surface-treated with an organosilicon compound such as an organopolysiloxane, organoalkoxysilane, organochlorosilane or a hexaorganodisilazane. The filler can be diatomaceous earth, finely crushed quartz, aluminum oxide, titanium oxide, iron oxide, cerium oxide, cerium hydroxide, magnesium oxide, zinc oxide, calcium carbonate, zirconium silicate, carbon black or ultramarine. A single filler or a combination of fillers can be used to reinforce the silicone polymer.

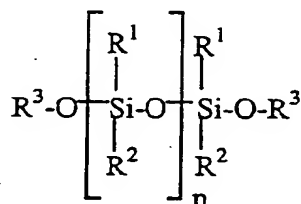
[0018] The amount of the filler can be in the range of from about 5 to about 200 parts by weight, desirably from about 10 to about 100 parts by weight and preferably from about 20 to about 60 parts by weight, per 100 parts by weight of silicone polymer.

[0019] The concentration of residual silanol groups on the surface of a filler can govern strength of hydrogen bonds between the silica and hydroxyl or oxygen groups in the silicone polymer chain. High concentrations of residual silanols in a filler cause "structuring" or "crepe hardening" of the final product in storage. This effect leads to difficulties in the processing of the material after it has been stored for extended periods. If the concentration of silanol functional groups in a filler is too high, a treating agent can be added to reduce the groups to a required concentration. The silanol reactant treating agent can react to reduce available groups to a concentration of between about 8 to about 2 hydroxyl groups/(nanometer)² of filler, preferably between about 5 to about 3 hydroxyl groups/(nanometer)² of filler. The surface-treated silica is a preferred filler in the invention, in an amount from about 10 to about 100 parts by weight, preferably from about 20 to about 60 parts by weight, per 100 parts by weight of silicone polymer.

[0020] In one embodiment, a treating agent is mixed into the filler along with the processing fluid or as the processing fluid to reduce filler silanol groups, to improve dispensability of the filler and/or to reduce the time required for aging of the silicon rubber, to prevent crepe hardening and/or to regulate plasticity. The treating agent can be the silanol-reacting reagent or another filler treating agent. The treating agent is preferably a silanol reactant treating agent when the filler is a silica or other silanol containing filler. The treating agent can be an organosilane, a low-viscosity polysiloxane or a silicone resin, which has a silanol group and/or an alkoxy group having 1 to 6 carbon atoms. Examples include

diphenyl-silanediol, dimethylsilanediol, methyltriethoxysilane and phenyltrimethoxysilane. The low-viscosity polysiloxane may contain one or more kinds of organic groups selected from a methyl group, a phenyl group, a vinyl group and a 3,3,3-trifluoropropyl group. The viscosity of the polysiloxane measured at 25°C is in the range of from about 1 to about 300 cP, preferably from about 5 to about 100 cP. Preferred silanol-reactant treating agents include silanol-stopped polydimethylsiloxane, octamethylcyclotetrasiloxane (D4) and hexamethyldisilazane (HMDZ).

[0021] The silicone polymer used in the compositions of the present invention is represented by recurring units of Formula I:



Formula I

wherein, R¹ independently at each occurrence represents C₁₋₄ alkyl, or C₂₋₄ alkylenes; R² independently at each occurrence represents C₁₋₄ alkyl, C₁₋₄ haloalkyl or C₂₋₄ alkylenes; R³ independently at each occurrence represents H, C₁₋₁₀ alkyl, C₂₋₄ alkylenes, C₄₋₆ cycloalkyl, OH or C₁₋₄ haloalkyl; and n represents an integer from 1,000 to 20,000.

[0022] A further preferred composition comprises a silicone polymer wherein, R¹ independently at each occurrence represents, CH₃ or CH=CH₂; R² independently at each occurrence represents, CH₃, CH=CH₂ or CH₂CH₂CF₃; R³ independently at each occurrence represents CH₃, CH=CH₂, OH or CH₂CH₂CF₃; and n represents an integer from about 4,000 to about 10,000.

[0023] Another embodiment provides a composition wherein the vinyl content of the silicone polymer ranges from about 0.05% to about 0.5 % by weight of the silicone polymer.

[0024] The silicone composition can also include other additives such as heat-resistance improvers such as oxides, hydroxides and fatty acid salts of metals, vulcanization reverse inhibitors, flame retardants such as platinum compounds, discoloration preventive agents, plasticizers such as silicone oil, internal release agent such as metal soaps, pigments and dyes.

[0025] During processing, an inert blanketing gas can be added to the compounding environment to suppress oxidative reactions between a flammable processing fluid such as HMDZ, and air. The amount of inert gas can be in the range of from about 20 to about 800 parts by weight, desirably from about 50 to about 600 parts by weight and preferably from about 100 to about 400 parts by weight per 100 parts by weight of the filler.

[0026] In an embodiment of the invention, fumed silica filler is mixed with a processing fluid that comprises HMDZ and water at a first location prior to addition of the silicone polymer. The amount of HMDZ can be in the range of from about 0.1 to about 100 parts by weight, desirably from about 0.5 to about 50 parts by weight and preferably from about 1.0 to about 20 parts by weight, per 100 parts by weight of the fumed silica. The amount of water can be in the range of from about 0.1 to about 100 parts by weight, desirably from about 0.5 to about 20 parts by weight and preferably from about 1 to about 10 parts by weight, per 100 parts by weight of the fumed silica.

[0027] The first extruder used in the invention can be a double screw extruder of the co-rotating intermeshing type. Or, the extruder can be a counter-rotating, non-intermeshing double screw extruder or a single screw extruder, either of the reciprocating or of the non-reciprocating type. The co-rotating, intermeshing double screw extruder is especially suited for the process of this invention due to its capability to produce the conditions needed for the preparation of the filler for compounding with the silicone polymer. Namely the extruder can provide a severe compounding and mixing energy. The extruder can have multiple addition ports for on-stream addition of the components of the compounded silicones.

[0028] The long extruder can be a counter-rotating, non-intermeshing double screw extruder, a co-rotating, intermeshing double screw extruder or a single screw reciprocating or non-reciprocating extruder. The long extruder provides the high rates of interfacial area generation that are needed to facilitate diffusion of volatile components through polymer-vapor interfaces. The volatile components are then diffused into a vapor space of the extruder and eliminated through vents. In one embodiment of the invention, the long extruder is a single screw reciprocating or non-reciprocating extruder of an L/D greater than 40.

[0029] Addition to the extruder of the individual components of the formulation, filler densification, chemical reaction between filler and treating agent, dispersion of filler into the silicone polymer to produce a homogeneous mixture and

devolatilization of the homogenized material are unit operations that are performed sequentially or concurrently. According to the invention, conditions in the extruder can be controlled to carry out these operations with complete product devolatilization. Hence in another embodiment, the invention relates to extrusion conditions, including compounding and devolatilization used to compound large amounts of filler particularly fumed silica and processing fluid, treating agent and silicone polymer.

[0030] Throughput and screw speed can be adjusted in the extruders to allow for efficient compounding and devolatilization. Low throughput under utilizes the capacity of manufacturing equipment. On the other hand, throughput is limited by the rate at which fumed silica can be added into an extruder. High screw speeds are needed for addition and dispersion of filler and dispersion of additives into the silicone matrix and for generation of surface area for devolatilization. However, temperature rises with viscous dissipation and screw speed. The use of severe screw speeds can result in thermal degradation of the silicone polymer. In the invention, scalable throughput with balanced mixing intensity provides effective compounding and reaction of silicone composition components with adequate process devolatilization due to material viscosity and extruder screw speed.

[0031] An extruder screw speed for either the first extruder or the long extruder can be between about 30 rpm and about 1000 rpm to provide a suitable balancing of mixing with frictional heat generation. Desirably, the screw speed is between about 100 rpm and about 800 rpm and preferably between about 300 rpm and about 500 rpm. A ratio of throughput to screw speed (lb/hour/rpm) for the first extruder or the long extruder can be between about 0.01 to about 500, desirably between 0.05 and about 200 and preferably between 1.0 and 100 (lb/hour/rpm).

[0032] Exterior first extruder barrel temperature can be between about 20°C and about 200°C, desirably between about 30°C and about 150°C, and preferably between about 40°C and about 100°C. Automatic tuning and programmable temperature controllers with time proportioning output in heating and cooling can be used for zone temperature control and thermocouples with plug ends can be used for barrel temperature measurements. For devolatilization, the long extruder barrel temperature can be between about 150°C and about 200°C, desirably between about 160°C and about 190°C and preferably between about 170°C and about 180°C.

[0033] The compositions produced by the process of the invention meet physical property standards for heat-vulcanizable silicone compositions. For example, the compositions can be characterized by a Williams Plasticity of greater than 100, Shore A hardness of greater than 20, tensile strength of greater than 750 psi, elongation at break of at least 100%, Tear B of at least 10 ppi, Specific Gravity of at least 1.05 and residual volatiles below 1 weight percent.

[0034] These and other features will become apparent from the following drawings and detailed discussion, which by way of example without limitation describe embodiments of the present invention.

[0035] In the drawings, FIGs. 1-3 are schematic representations of silicone compounding processes and systems for carrying out embodiments of the invention. In FIG. 1, an overall process and system for compounding filled silicone compositions is designated 10. The system includes first extruder 12 and long extruder 14.

[0036] The first extruder 12 can be a double screw extruder of the co-rotating or counter-rotating, intermeshing type, a counter-rotating, non-intermeshing type, a single-screw reciprocating or a single screw non-reciprocating type. Multiple extruder stages can be used as the first extruder to compound filler, treating agent and silicone polymer. The homogenous material can be fed to the long extruder. The compounded material can be force-fed from the first extruder to the long extruder by the use of pressure generated internally by the rotation of the extruder screws.

[0037] The long extruder 14 can be a counter-rotating, non-intermeshing double screw extruder, a co-rotating, intermeshing double screw extruder or a single screw reciprocating or non-reciprocating extruder of an L/D of greater than 50. The long extruder 14 is designed to operate at high temperatures and to provide the open surface areas necessary for devolatilization. The long extruder 14 provides the high pressurization necessary to pump compounded product out of the final extruder 14.

[0038] In the process of FIG. 1, filler, processing fluid, which in this embodiment includes a treating agent and silicone polymer are continuously supplied from respective storage tanks 16, 18 and 20 by means of respective feed pumps 22, 24 and 26 into first extruder 12. In extruder 12, the filler, processing fluid and silicone polymer are continuously compounded and discharged as an extrudate, which is force fed 28 into long extruder 14 having a L/D ratio of greater than 50. Long extruder 14 is provided with vent 30 for extrudate devolatilization. Vent 30 can represent a plurality of vents in applications that require more than one vent for effective devolatilization. After venting, a compounded and devolatilized filled silicone composition is discharged through long extruder end 32.

[0039] In FIG. 2, the system 40 includes mixer 42, first extruder 44 and long extruder 46. In the process of FIG. 2, filler and silicone polymer are combined and mixed in mixer 42 to form a premix which is fed 48 into first extruder 44. Additional filler, processing fluid and additional silicone polymer are continuously supplied from respective storage tanks 50, 52 and 54 by means of respective feed pumps 56, 58 and 60 into first extruder 44 where they are continuously compounded and discharged as an extrudate, which is continuously force fed 62 into long extruder 46 having a L/D ratio of greater than 50. Long extruder 46 is provided with vent 64 for extrudate devolatilization. After venting, a compounded and devolatilized filled silicone composition is discharged through long extruder end 66.

[0040] In FIG. 3, the system 70 includes reactor 72, mixer 74, first extruder 76 and long extruder 78. A raw filler and

EP 1 110 691 A2

a treating agent are reacted in reactor 72 to form a pretreated filler, which is discharged 80 to mixer 74. Silicone polymer is added 82 to the pretreated filler and mixed in mixer 74 to form a premix. The premix is fed 84 into first extruder 76. Additional filler, processing fluid and silicone polymer are continuously supplied from respective storage tanks 86, 88 and 90 by means of respective feed pumps 92, 94 and 96 into first extruder 76 where they are continuously compounded and force fed 98 as an extrudate into long extruder 78 having a L/D ratio of greater than 50. Long extruder 78 is provided with vent 100 for extrudate devolatilization. After venting, a compounded and devolatilized filled silicone composition is discharged through long extruder end 102.

[0041] Examples

[0042] In these Examples, compounding was performed in a 2-inch diameter, co-rotating, intermeshing double screw first extruder. The extruder had a length to diameter ratio of about 8. The barrel of this extruder could be heated or cooled but had no venting capability. The first extruder screws were operated at between 50 and 400 rpm. The first extruder was capable of compounding filler and silicone polymer compositions into an homogeneous filled silicone composition.

[0043] Devolatilization was performed in a 30 mm diameter, counter-rotating, non-intermeshing double screw long extruder. The long extruder had a length-to-diameter ratio of 60. The long extruder was equipped with three vacuum vents (designated upstream, downstream and intermediate) vents for devolatilization. The vents were capable of being operated at absolute pressures as low as a few millimeters of mercury. The long extruder had segmented barrels that could be maintained at a constant temperature using recirculating heated oil. The long extruder included two 1.05 inch outside diameter (OD) cylindrical elements before the upstream vent, two 0.95 inch OD cylindrical screw elements between the upstream and the intermediate vent, two 0.9 inch OD cylindrical screw elements between the intermediate and downstream vent and a last 0.915 inch single, forward flighted screw element that could be used for pumping. The screws were capable of rotation at speeds between 300 and 500 rpm.

[0044] In the processes described in the Examples, all of the components of the compositions were added to the co-rotating, intermeshing first extruder at an upstream portion of the process. A particulate concentrate of D4 treated fumed silica and high molecular weight silica polymer was prepared in a Henschel batch mixer and was added at the first extruder feed throat using an Acrison feeder. A mixture of liquid additives was incorporated at the first upstream injection port and silicone polymer was added at a side feeding port by means of a positive displacement gear pump.

[0045] Table 1 includes operating conditions and feed rates at the first co-rotating, intermeshing extruder for each of runs 1001 through 1004.

Table 1

	1001	1002	1003	1004
screw speed (rpm)	100	100	250	250
filler/polymer premix (lb/hr)	39.985	31.984	32.051	39.936
silicone polymer (lb/hr)	18.17	26.09	26.09	18.17
vinyl silazane additive (lb/hr)	0.024	0.024	0.066	0.066
vinyl-stopped silicone additive (lb/hr)	0.547	1.251	0.547	1.251
silanol stopped silicone additive (lb/hr)	0.541	1.260	0.541	1.260

[0046] Table 2 includes conditions for devolatilization in the long extruder.

Table 2

	1001	1002	1003	1004
screw speed (rpm)	400	375	375	450
pump vacuum (In Hg)	29	29	29	29
vent pressure (mm Hg)	2	2	3.5	3.7
Number of Vents	2	2	2	2

[0047] Table 3 shows product properties for the Examples of Tables 1 and 2.

Table 3

	1001	1002	1003	1004
Shore A Hardness	51.3	41.8	38.0	51.1
Tensile Strength (ppi)	1206	876	1077	1063
Elongation (%)	404	456	516	420
Tear B (ppi)	93	68	72	147
Residual Volatiles (weight percent)	0.61	0.72	0.65	0.77
Specific Gravity	1.132	1.094	1.097	1.129

[0048] The Examples illustrate the invention conducted with ranges of formulations and process conditions. The results of Table 3 show that the process of the invention can produce materials within physical property standards for filled heat-vulcanizable silicone compositions.

Claims

1. A process of compounding a filled silicone composition, comprising:

compounding a filler, processing fluid and silicone polymer in a first compounding apparatus (12) to produce a dispersed composition containing volatiles; and

devolatilizing said dispersed composition in a long extruder (14) having an L/D ratio of greater than 50 to devolatilize said dispersed composition.

2. The process of claim 1, wherein said filler is a raw, untreated silica.

3. The process of claim 1, wherein said filler is a pretreated filler with treating agent.

4. The process of claim 3, wherein said treating agent is a silanol-reacting treating agent.

5. The process of any preceding claim, wherein said compounding apparatus (12) comprises a first co-rotating, intermeshing double screw extruder; and said long extruder (14) comprises a counter-rotating, non-intermeshing double screw extruder; a co-rotating; non-intermeshing double screw extruder; or a single screw reciprocating extruder or single screw non-reciprocating extruder.

6. The process of any one of claims 1 to 4, wherein said compounding apparatus (12) comprises a single screw reciprocating extruder or a single screw non-reciprocating extruder; and said long extruder (14) comprises a co-rotating, intermeshing double screw extruder, a counter-rotating; intermeshing double screw extruder or a counter-rotating, non-intermeshing double screw extruder.

7. The process of claim 1, wherein said filler contains silanol groups and said processing fluid is a treating agent comprising silanol-stopped polydimethylsiloxane, octamethylcyclotetrasiloxane (D4) or hexamethyldisilazane (HMDZ).

8. The process of claim 1, wherein said processing fluid is selected from the group consisting of silanol-stopped polydimethylsiloxane, vinyl-stopped dimethyl-methylvinylsiloxane and hydroxy-terminated polydimethyl-methylvinylsiloxane

9. The process of any preceding claim, wherein said compounding is completed in a compounding apparatus (12) of at least some length that is about 30 diameters or less in length.

10. The process of any preceding claim, comprising controlling said compounding to provide a total throughput to screw speed ratio in said compounding apparatus (12) between about 0.01 and about 500 (lb/hour/rpm) and con-

trolling said devolatilization to provide a total throughput to screw speed ratio in said extruder (14) between about 0.01 and about 500 (lb/hour/rpm).

5 11. The process of any preceding claim, comprising controlling said compounding to provide a total throughput to screw speed ratio in said compounding apparatus (12) between about 0.05 and about 200 (lb/hour/rpm) and controlling said devolatilization to provide a total throughput to screw speed ratio in said extruder (14) between about 0.05 and about 200 (lb/hour/rpm).

10 12. The process of any preceding claim, comprising controlling said compounding to provide a total throughput to screw speed ratio in said compounding apparatus (12) between about 1.0 and about 100 (lb/hour/rpm) and controlling said devolatilization to provide a total throughput to screw speed ratio in said extruder (14) between about 1.0 and about 100 (lb/hour/rpm).

15 13. A process of compounding a filled silicone composition, comprising:

forming a premix of filler and silicone polymer in a first mixer;

20 compounding said premix with further filler and silicone polymer and added processing fluid in a first compounding apparatus (12) to produce a dispersed composition containing volatiles; and

devolatilizing said dispersed composition in a long extruder (14) having an L/D ratio of greater than 50 to devolatilize said dispersed composition.

25 14. A system (10) for compounding filled silicone compositions, comprising:

a first compounding apparatus (12);

and a long extruder (14) having an L/D ratio of greater than 50, connected to said first compounding apparatus (12) to continuously receive a dispersed composition containing volatiles from said first compounding apparatus (12).

30 15. A system (40) for compounding filled silicone compositions, comprising:

a mixing apparatus (42);

35 a first compounding apparatus (44);

and a long extruder (46) having an L/D ratio of greater than 50, connected to said first compounding apparatus (44) to continuously receive a dispersed composition containing volatiles from said first compounding apparatus (44).

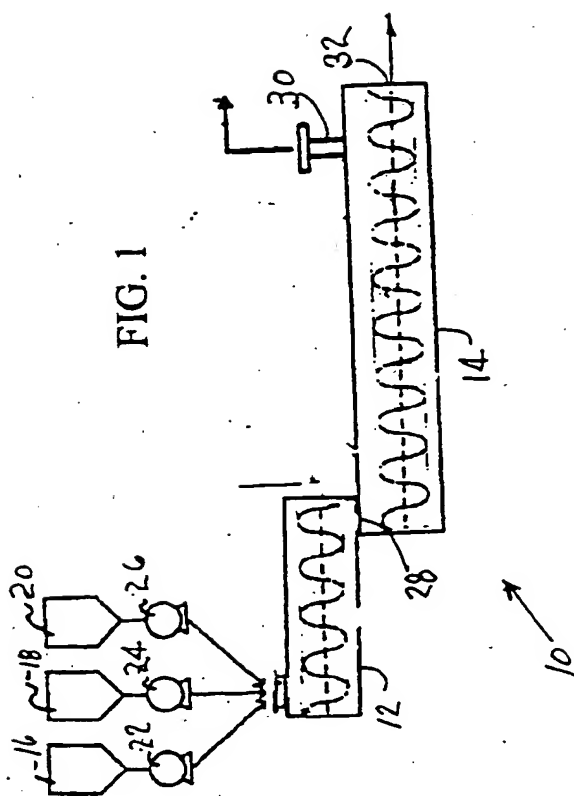


FIG. 2

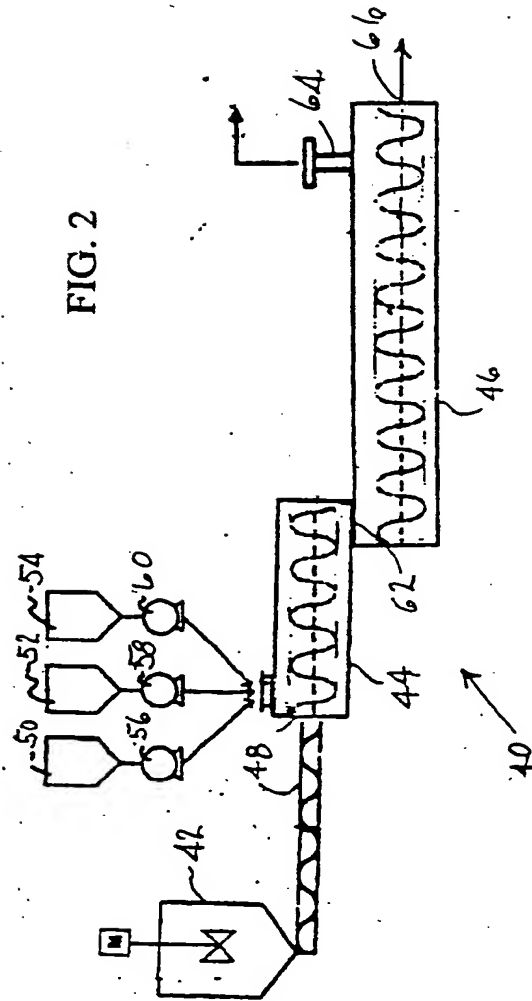
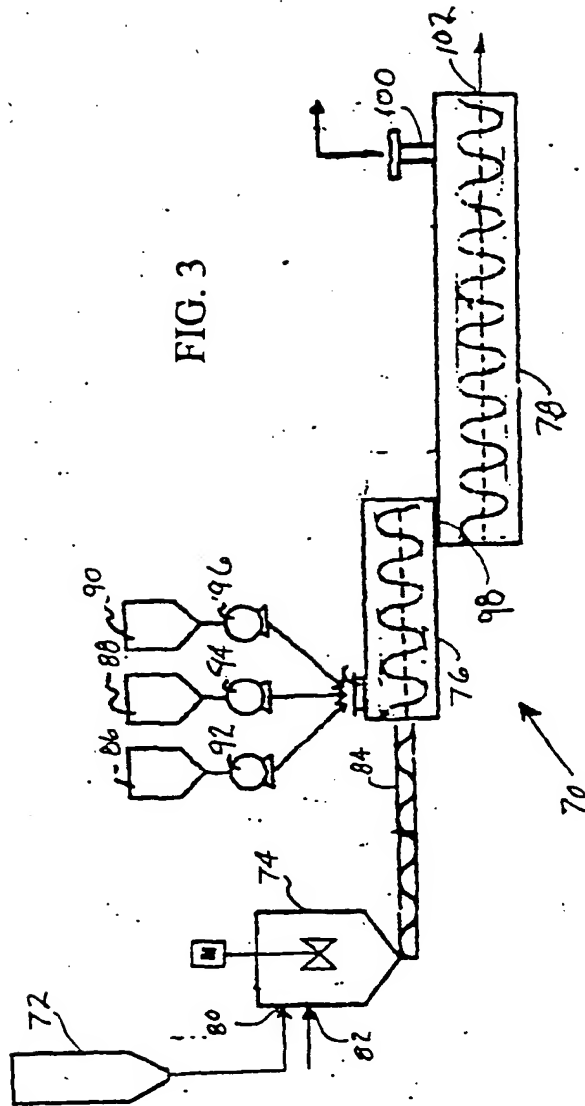


FIG. 3



(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 110 691 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
20.03.2002 Bulletin 2002/12

(51) Int Cl.7: **B29C 47/50**
// B29K83:00

(43) Date of publication A2:
27.06.2001 Bulletin 2001/26

(21) Application number: 00310967.5

(22) Date of filing: 08.12.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 21.12.1999 US 468622

(71) Applicant: GENERAL ELECTRIC COMPANY
Schenectady, NY 12345 (US)

(72) Inventors:
• Silvi, Norberto
Clifton Park, New York 12065 (US)
• Dean, Robert Edward
Rexford, New York 12148 (US)

- Giammattei, Mark Howard
Selkirk, New York 12158 (US)
- Carbone, John William
Schenectady, New York 12309 (US)
- Singh, Navjot
Clifton Park, New York 12065 (US)
- Rocha, Gerardo
Waterford, New York 12188 (US)
- Banevicius, John Peter
Clifton Park, New York 12065 (US)

(74) Representative: Szary, Anne Catherine, Dr. et al
London Patent Operation,
GE International, Inc.,
Essex House,
12-13 Essex Street
London WC2R 3AA (GB)

(54) Compounding filled silicone compositions

(57) High levels of inorganic filler, processing fluid and silicone polymer are compounded and devolatilized into homogeneous filled compositions with requisite reinforcing properties and levels of volatiles. In the method, a filler, processing fluid and silicone polymer are

mixed in a first compounding apparatus (12,44,76) to produce a dispersed composition containing volatiles. The dispersed composition is then compounded in a long extruder (14,46,78) having an L/D ratio of greater than 50 to devolatilize the dispersed composition.

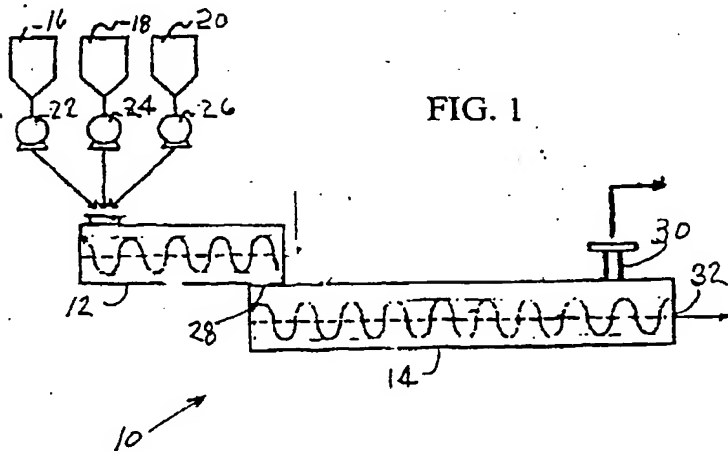


FIG. 1

EP 1 110 691 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 31 0967

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	EP 0 258 159 A (RHONE POULENC CHIMIE) 2 March 1988 (1988-03-02) * column 4, line 34 - line 37 * * column 9, line 35 - column 10, line 27; figures 1,2 *	1-4,7-9, 14	B29C47/50 //B29K83:00
A	GB 1 174 219 A (WERNER & PFLEIDERER) 17 December 1969 (1969-12-17) * the whole document *	1-15	
A	EP 0 622 398 A (DOW CORNING TORAY SILICONE) 2 November 1994 (1994-11-02) * the whole document *	1-15	
A	EP 0 568 891 A (BAYER AG) 10 November 1993 (1993-11-10) * page 3, line 3 - line 54; figure 1 *	1-15	
A	WO 92 13694 A (WACKER CHEMIE GMBH) 20 August 1992 (1992-08-20) * page 10, paragraph 2 - page 12, paragraph 3; claims 1-5 *	1-15	
A	EP 0 512 730 A (DOW CORNING SA) 11 November 1992 (1992-11-11) * column 3, line 54 - column 6, line 26 *	1-15	B29B B29C B01F
D,A	US 5 198 171 A (WATANABE JUNICHIRO ET AL) 30 March 1993 (1993-03-30) * the whole document *	1-15	
A	EP 0 462 032 A (RHONE POULENC CHIMIE) 18 December 1991 (1991-12-18) * page 4, line 17 - line 18; figure 1; example 1 *	1-15	
-/-			
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 29 January 2002	Examiner Van Nieuwenhuize, O
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons</p> <p>& : member of the same patent family, corresponding document</p>			

EPO FORM 1503 03/82 (PO/C01)



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 31 0967

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
A	WO 81 00259 A (BUSS AG ;IROQUOIS CHEM GMBH (CH); STOJANOVIC I (CH); SCHIMANSKI P) 5 February 1981 (1981-02-05) * claim 2; figure 1 *	14, 15	
A	US 4 063 718 A (KOCH KLAUS) 20 December 1977 (1977-12-20) * claim 1 *	14	
A	EP 0 776 751 A (DOW CORNING TORAY SILICONE) 4 June 1997 (1997-06-04) * page 2, line 49; claim 1; figure 2 *	1	
			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 29 January 2002	Examiner Van Nieuwenhuize, O
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 1603 03 82 (P01C01)

ANNEX TO THE EUROPEAN SEARCH REPORT ON EUROPEAN PATENT APPLICATION NO.

EP 00 31 0967

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

29-01-2002

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0258159	A	02-03-1988	FR 2602710 A1	19-02-1988
			BR 8704547 A	26-04-1988
			DE 3762023 D1	03-05-1990
			EP 0258159 A1	02-03-1988
GB 1174219	A	17-12-1969	DE 1719321 A1	27-07-1972
			FR 1583057 A	17-10-1969
			NL 6806477 A	11-11-1968
EP 0622398	A	02-11-1994	JP 6313046 A	08-11-1994
			DE 69410525 D1	02-07-1998
			DE 69410525 T2	15-10-1998
			EP 0622398 A1	02-11-1994
			US 5409978 A	25-04-1995
EP 0568891	A	10-11-1993	DE 4215205 C1	05-01-1994
			AT 177459 T	15-03-1999
			CA 2095572 A1	09-11-1993
			DE 59309420 D1	15-04-1999
			EP 0568891 A1	10-11-1993
			ES 2131082 T3	16-07-1999
			JP 6032909 A	08-02-1994
			KR 256932 B1	15-05-2000
WO 9213694	A	20-08-1992	DE 4103602 A1	13-08-1992
			AT 115903 T	15-01-1995
			CS 9200034 A3	12-08-1992
			DE 59201026 D1	02-02-1995
			WO 9213694 A1	20-08-1992
			EP 0570387 A1	24-11-1993
			ES 2065774 T3	16-02-1995
			JP 2531912 B2	04-09-1996
			JP 6503048 T	07-04-1994
			KR 9700921 B1	21-01-1997
			RU 2091222 C1	27-09-1997
			US 6124392 A	26-09-2000
EP 0512730	A	11-11-1992	DE 69208387 D1	28-03-1996
			DE 69208387 T2	25-07-1996
			EP 0512730 A2	11-11-1992
			JP 5131516 A	28-05-1993
			US 5227111 A	13-07-1993
US 5198171	A	30-03-1993	JP 2102007 A	13-04-1990
			JP 8002532 B	17-01-1996
			KR 9303375 B1	26-04-1993

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 31 0967

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

29-01-2002

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
EP 0462032	A	18-12-1991	FR	2663340 A1	20-12-1991
			AP	221 A	21-10-1992
			AT	133188 T	15-02-1996
			AU	7625691 A	19-12-1991
			CA	2044199 A1	14-12-1991
			DE	69116443 D1	29-02-1996
			DE	69116443 T2	22-08-1996
			DK	462032 T3	12-02-1996
			EP	0462032 A1	18-12-1991
			ES	2082169 T3	16-03-1996
			GR	3019368 T3	30-06-1996
			JP	4226157 A	14-08-1992
			JP	7037062 B	26-04-1995
			US	5531923 A	02-07-1996
WO 8100259	A	05-02-1981	BE	884296 A1	03-11-1980
			WO	8100259 A1	05-02-1981
			DE	3049740 D2	11-02-1982
			EP	0032142 A1	22-07-1981
			IT	1132527 B	02-07-1986
US 4063718	A	20-12-1977	DE	2543328 A1	31-03-1977
			CH	613150 A5	14-09-1979
			DD	126909 A5	24-08-1977
			FR	2325493 A1	22-04-1977
			GB	1567026 A	08-05-1980
			IT	1072445 B	10-04-1985
			JP	52042554 A	02-04-1977
			SE	7610435 A	30-03-1977
EP 0776751	A	04-06-1997	JP	9155950 A	17-06-1997
			DE	69611815 D1	29-03-2001
			DE	69611815 T2	27-09-2001
			EP	0776751 A1	04-06-1997
			US	5721307 A	24-02-1998

EPO FORM P4459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

THIS PAGE BLANK (USPTO)